



UHID	12051116	Date	08/10/2022		
Name	Ms. Pimki Kumari Rajak	Sex	Female	Age	40
OPD	Pap Smear 20	Health Check-up			

Drug allergy:
 Sys illness:

40 yrs / Polo

Single / separated

LMP: 20-9-22

PMC: 3/30d, RMP

Rsp - cop / ng / pap

Breast examⁿ @

Adv

- Self breast examⁿ monthly
- Pap smear - yearly
- mammography
 us & Pelvis } yearly

heha



UHID	12051116	Date	08/10/2022		
Name	Ms. Pimki Kumari Rajak	Sex	Female	Age	40
OPD	Ophthal 14	Health Check-up			

Obs. No.

Drug allergy: → Not known
 Sys illness: → NO.

Mes Thyroid (since 2 weeks).

Visual acuity → RE 6/gp
 → L 6/gp

Refraction → RE Plano / -1.00 x 90° 6/g
 → L Plano / -1.00 x 90° 6/g
 Add →

I.O.P. → RG → 16.2
 → LG → 17.1

Shilpa



UHID	12051116	Date	08/10/2022		
Name	Ms. Pimki Kumari Rajak	Sex	Female	Age	40
OPD	Dental 12	Health Check-up			

Drug allergy:
Sys illness:

1) Stain+

Calculus+++

Adv

1) Oral propylaxir

BHI

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CLIENT PATIENT ID : UID:12051116

ACCESSION NO : **0022VJ001489** AGE : 40 Years SEX : Female DATE OF BIRTH : 17/06/1982

DRAWN : 08/10/2022 10:10 RECEIVED : 08/10/2022 10:10 REPORTED : 08/10/2022 14:59

CLIENT NAME : **FORTIS VASHI-CHC -SPLZD**

REFERRING DOCTOR : SELF

CLINICAL INFORMATION :

UID:12051116 REQNO-1305061

CORP-OPD

BILLNO-1501220PCR050212

BILLNO-1501220PCR050212

Test Report Status	Final	Results	Biological Reference Interval	Units
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KIDNEY PANEL - 1**SERUM BLOOD UREA NITROGEN**

BLOOD UREA NITROGEN 9 6 - 20 mg/dL

METHOD : UREASE - UV

CREATININE EGFR- EPI

CREATININE 0.73 0.60 - 1.10 mg/dL

METHOD : ALKALINE PICRATE KINETIC JAFFES

AGE 40 years

GLOMERULAR FILTRATION RATE (FEMALE) 106.55 Refer Interpretation Below mL/min/1.73m2

METHOD : CALCULATED PARAMETER

BUN/CREAT RATIO

BUN/CREAT RATIO 12.33 5.00 - 15.00

METHOD : CALCULATED PARAMETER

URIC ACID, SERUM

URIC ACID 3.2 2.6 - 6.0 mg/dL

METHOD : URICASE UV

TOTAL PROTEIN, SERUM

TOTAL PROTEIN 7.3 6.4 - 8.2 g/dL

METHOD : BIURET

ALBUMIN, SERUM

ALBUMIN 3.5 3.4 - 5.0 g/dL

METHOD : BCP DYE BINDING

GLOBULIN

GLOBULIN 3.8 2.0 - 4.1 g/dL

METHOD : CALCULATED PARAMETER

ELECTROLYTES (NA/K/CL), SERUM

SODIUM 139 136 - 145 mmol/L

METHOD : ISE INDIRECT

POTASSIUM 4.31 3.50 - 5.10 mmol/L

METHOD : ISE INDIRECT

CHLORIDE 105 98 - 107 mmol/L

METHOD : ISE INDIRECT

Interpretation(s)

SERUM BLOOD UREA NITROGEN-

Causes of Increased levels

Pre renal

- High protein diet, Increased protein catabolism, GI haemorrhage, Cortisol, Dehydration, CHF Renal

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- Renal Failure
- Post Renal
- Malignancy, Nephrolithiasis, Prostatism

Causes of decreased levels

- Liver disease
- SIADH.

CREATININE EGFR- EPI-

GFR— Glomerular filtration rate (GFR) is a measure of the function of the kidneys. The GFR is a calculation based on a serum creatinine test. Creatinine is a muscle waste product that is filtered from the blood by the kidneys and excreted into urine at a relatively steady rate. When kidney function decreases, less creatinine is excreted and concentrations increase in the blood. With the creatinine test, a reasonable estimate of the actual GFR can be determined.

A GFR of 60 or higher is in the normal range.

A GFR below 60 may mean kidney disease.

A GFR of 15 or lower may mean kidney failure.

Estimated GFR (eGFR) is the preferred method for identifying people with chronic kidney disease (CKD). In adults, eGFR calculated using the Modification of Diet in Renal Disease (MDRD) Study equation provides a more clinically useful measure of kidney function than serum creatinine alone.

The CKD-EPI creatinine equation is based on the same four variables as the MDRD Study equation, but uses a 2-slope spline to model the relationship between estimated GFR and serum creatinine, and a different relationship for age, sex and race. The equation was reported to perform better and with less bias than the MDRD Study equation, especially in patients with higher GFR. This results in reduced misclassification of CKD.

The CKD-EPI creatinine equation has not been validated in children & will only be reported for patients = 18 years of age. For pediatric and childrens, Schwartz Pediatric Bedside eGFR (2009) formulae is used. This revised "bedside" pediatric eGFR requires only serum creatinine and height.

URIC ACID, SERUM-

Causes of Increased levels

Dietary

- High Protein Intake.
- Prolonged Fasting,
- Rapid weight loss.

Gout

Lesch nyhan syndrome.

Type 2 DM.

Metabolic syndrome.

Causes of decreased levels

- Low Zinc Intake
- OCP's
- Multiple Sclerosis

Nutritional tips to manage increased Uric acid levels

- Drink plenty of fluids
- Limit animal proteins
- High Fibre foods
- Vit C Intake
- Antioxidant rich foods

TOTAL PROTEIN, SERUM-

Serum total protein,also known as total protein, is a biochemical test for measuring the total amount of protein in serum..Protein in the plasma is made up of albumin and globulin

Higher-than-normal levels may be due to: Chronic inflammation or infection, including HIV and hepatitis B or C, Multiple myeloma, Waldenström's disease
Lower-than-normal levels may be due to: Agammaglobulinemia, Bleeding (hemorrhage), Burns, Glomerulonephritis, Liver disease, Malabsorption, Malnutrition, Nephrotic syndrome, Protein-losing enteropathy etc.

ALBUMIN, SERUM-

Human serum albumin is the most abundant protein in human blood plasma. It is produced in the liver. Albumin constitutes about half of the blood serum protein. Low blood albumin levels (hypoalbuminemia) can be caused by: Liver disease like cirrhosis of the liver, nephrotic syndrome, protein-losing enteropathy, Burns, hemodilution, increased vascular permeability or decreased lymphatic clearance, malnutrition and wasting etc.

ELECTROLYTES (NA/K/CL), SERUM-

Sodium levels are Increased in dehydration, cushing's syndrome, aldosteronism & decreased in Addison's disease, hypopituitarism, liver disease. Hypokalemia (low K) is common in vomiting, diarrhea, alcoholism, folic acid deficiency and primary aldosteronism. Hyperkalemia may be seen in end-stage renal failure, hemolysis, trauma, Addison's disease, metabolic acidosis, acute starvation, dehydration, and with rapid K infusion. Chloride is increased in dehydration, renal tubular acidosis (hyperchloremia metabolic acidosis), acute renal failure, metabolic acidosis associated with prolonged diarrhea and loss of sodium bicarbonate, diabetes insipidus, adrenocortical hyperfunction, salicylate intoxication and with excessive infusion of isotonic saline or extremely high dietary intake of salt. Chloride is decreased in overhydration, chronic respiratory acidosis, salt-losing nephritis, metabolic alkalosis, congestive heart failure, Addisonian crisis, certain types of metabolic acidosis, persistent gastric secretion and prolonged vomiting.

HAEMATOLOGY**SRL Ltd**

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ERYTHRO SEDIMENTATION RATE, BLOOD

SEDIMENTATION RATE (ESR) 16 0 - 20 mm at 1 hr
METHOD : WESTERGREN METHOD

CBC-5, EDTA WHOLE BLOOD**BLOOD COUNTS, EDTA WHOLE BLOOD**

HEMOGLOBIN 12.0 12.0 - 15.0 g/dL
METHOD : SPECTROPHOTOMETRY
RED BLOOD CELL COUNT 4.16 3.8 - 4.8 mil/ μ L
METHOD : ELECTRICAL IMPEDANCE
WHITE BLOOD CELL COUNT 6.88 4.0 - 10.0 thou/ μ L
METHOD : DOUBLE HYDRODYNAMIC SEQUENTIAL SYSTEM(DHSS)CYTOMETRY
PLATELET COUNT 234 150 - 410 thou/ μ L
METHOD : ELECTRICAL IMPEDANCE

RBC AND PLATELET INDICES

HEMATOCRIT **34.7** **Low** 36 - 46 %
METHOD : CALCULATED PARAMETER
MEAN CORPUSCULAR VOLUME 83.3 83 - 101 fL
METHOD : CALCULATED PARAMETER
MEAN CORPUSCULAR HEMOGLOBIN 28.8 27.0 - 32.0 pg
METHOD : CALCULATED PARAMETER
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION 34.5 31.5 - 34.5 g/dL
METHOD : CALCULATED PARAMETER
MENTZER INDEX 20.0
RED CELL DISTRIBUTION WIDTH **16.1** **High** 11.6 - 14.0 %
METHOD : CALCULATED PARAMETER
MEAN PLATELET VOLUME **11.9** **High** 6.8 - 10.9 fL
METHOD : CALCULATED PARAMETER

WBC DIFFERENTIAL COUNT - NLR

NEUTROPHILS 64 40 - 80 %
METHOD : FLOW CYTOMETRY
ABSOLUTE NEUTROPHIL COUNT 4.40 2.0 - 7.0 thou/ μ L
METHOD : CALCULATED PARAMETER
LYMPHOCYTES 27 20 - 40 %
METHOD : FLOW CYTOMETRY

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ABSOLUTE LYMPHOCYTE COUNT		1.86	1.0 - 3.0	thou/ μ L
METHOD : CALCULATED PARAMETER				
NEUTROPHIL LYMPHOCYTE RATIO (NLR)		2.3		
METHOD : CALCULATED PARAMETER				
EOSINOPHILS		4	1 - 6	%
METHOD : FLOW CYTOMETRY				
ABSOLUTE EOSINOPHIL COUNT		0.28	0.02 - 0.50	thou/ μ L
METHOD : CALCULATED PARAMETER				
MONOCYTES		5	2 - 10	%
METHOD : FLOW CYTOMETRY				
ABSOLUTE MONOCYTE COUNT		0.34	0.2 - 1.0	thou/ μ L
METHOD : CALCULATED PARAMETER				
BASOPHILS		0	0 - 2	%
METHOD : FLOW CYTOMETRY				
ABSOLUTE BASOPHIL COUNT		0	Low 0.02 - 0.10	thou/ μ L
METHOD : CALCULATED PARAMETER				
DIFFERENTIAL COUNT PERFORMED ON:		EDTA SMEAR		
MORPHOLOGY				
RBC		PREDOMINANTLY NORMOCYTIC NORMOCHROMIC		
METHOD : MICROSCOPIC EXAMINATION				
WBC		NORMAL MORPHOLOGY		
METHOD : MICROSCOPIC EXAMINATION				
PLATELETS		ADEQUATE		
METHOD : MICROSCOPIC EXAMINATION				

Interpretation(s)**ERYTHRO SEDIMENTATION RATE, BLOOD-**

Erythrocyte sedimentation rate (ESR) is a non-specific phenomena and is clinically useful in the diagnosis and monitoring of disorders associated with an increased production of acute phase reactants. The ESR is increased in pregnancy from about the 3rd month and returns to normal by the 4th week post partum. ESR is influenced by age, sex, menstrual cycle and drugs (eg. corticosteroids, contraceptives). It is especially low (0 -1mm) in polycythaemia, hypofibrinogenemia or congestive cardiac failure and when there are abnormalities of the red cells such as poikilocytosis, spherocytosis or sickle cells.

Reference :

1. Nathan and Oski's Haematology of Infancy and Childhood, 5th edition
2. Paediatric reference intervals. AACC Press, 7th edition. Edited by S. Soldin
3. The reference for the adult reference range is "Practical Haematology by Dacie and Lewis, 10th Edition"

RBC AND PLATELET INDICES-

Mentzer index (MCV/RBC) is an automated cell-counter based calculated screen tool to differentiate cases of Iron deficiency anaemia(>13) from Beta thalassaemia trait (<13) in patients with microcytic anaemia. This needs to be interpreted in line with clinical correlation and suspicion. Estimation of HbA2 remains the gold standard for diagnosing a case of beta thalassaemia trait.

WBC DIFFERENTIAL COUNT - NLR-The optimal threshold of 3.3 for NLR showed a prognostic possibility of clinical symptoms to change from mild to severe in COVID positive patients. When age = 49.5 years old and NLR = 3.3, 46.1% COVID-19 patients with mild disease might become severe. By contrast, when age < 49.5 years old and NLR < 3.3, COVID-19 patients tend to show mild disease.

(Reference to - The diagnostic and predictive role of NLR, d-NLR and PLR in COVID-19 patients ; A.-P. Yang, et al.; International Immunopharmacology 84 (2020) 106504 This ratio element is a calculated parameter and out of NABL scope.

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IMMUNOHAEMATOLOGY**ABO GROUP & RH TYPE, EDTA WHOLE BLOOD**

ABO GROUP TYPE B

METHOD : TUBE AGGLUTINATION

RH TYPE NEGATIVE

METHOD : TUBE AGGLUTINATION

Interpretation(s)

ABO GROUP & RH TYPE, EDTA WHOLE BLOOD-

Blood group is identified by antigens and antibodies present in the blood. Antigens are protein molecules found on the surface of red blood cells. Antibodies are found in plasma. To determine blood group, red cells are mixed with different antibody solutions to give A,B,O or AB.

Disclaimer: "Please note, as the results of previous ABO and Rh group (Blood Group) for pregnant women are not available, please check with the patient records for availability of the same."

The test is performed by both forward as well as reverse grouping methods.

BIO CHEMISTRY**LIVER FUNCTION PROFILE, SERUM**

BILIRUBIN, TOTAL	0.30	0.2 - 1.0	mg/dL
METHOD : JENDRASSIK AND GROFF			
BILIRUBIN, DIRECT	0.13	0.0 - 0.2	mg/dL
METHOD : JENDRASSIK AND GROFF			
BILIRUBIN, INDIRECT	0.17	0.1 - 1.0	mg/dL
METHOD : CALCULATED PARAMETER			
TOTAL PROTEIN	7.3	6.4 - 8.2	g/dL
METHOD : BIURET			
ALBUMIN	3.5	3.4 - 5.0	g/dL
METHOD : BCP DYE BINDING			
GLOBULIN	3.8	2.0 - 4.1	g/dL
METHOD : CALCULATED PARAMETER			
ALBUMIN/GLOBULIN RATIO	0.9	Low 1.0 - 2.1	RATIO
METHOD : CALCULATED PARAMETER			
ASPARTATE AMINOTRANSFERASE (AST/SGOT)	20	15 - 37	U/L
METHOD : UV WITH P5P			
ALANINE AMINOTRANSFERASE (ALT/SGPT)	40	High < 34.0	U/L
METHOD : UV WITH P5P			
ALKALINE PHOSPHATASE	71	30 - 120	U/L

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METHOD : PNPP-ANP			
GAMMA GLUTAMYL TRANSFERASE (GGT)		17	5 - 55 U/L
METHOD : GAMMA GLUTAMYL CARBOXY 4NITROANILIDE			
LACTATE DEHYDROGENASE		122	100 - 190 U/L
METHOD : LACTATE -PYRUVATE			
GLUCOSE, FASTING, PLASMA			
GLUCOSE, FASTING, PLASMA		82	74 - 99 mg/dL
METHOD : HEXOKINASE			
GLYCOSYLATED HEMOGLOBIN, EDTA WHOLE BLOOD			
GLYCOSYLATED HEMOGLOBIN (HBA1C)		5.4	Non-diabetic: < 5.7 Pre-diabetics: 5.7 - 6.4 Diabetics: > or = 6.5 ADA Target: 7.0 Action suggested: > 8.0 %
METHOD : HB VARIANT (HPLC)			
MEAN PLASMA GLUCOSE		108.3	< 116.0 mg/dL
METHOD : CALCULATED PARAMETER			
CORONARY RISK PROFILE (LIPID PROFILE), SERUM			
CHOLESTEROL		183	< 200 Desirable 200 - 239 Borderline High >= 240 High mg/dL
METHOD : ENZYMATIC/COLORIMETRIC, CHOLESTEROL OXIDASE, ESTERASE, PEROXIDASE			
TRIGLYCERIDES		118	< 150 Normal 150 - 199 Borderline High 200 - 499 High >=500 Very High mg/dL
METHOD : ENZYMATIC ASSAY			
HDL CHOLESTEROL		56	< 40 Low >=60 High mg/dL
METHOD : DIRECT MEASURE - PEG			
DIRECT LDL CHOLESTEROL		105	< 100 Optimal 100 - 129 Near or above optimal 130 - 159 Borderline High 160 - 189 High >= 190 Very High mg/dL
METHOD : DIRECT MEASURE WITHOUT SAMPLE PRETREATMENT			
NON HDL CHOLESTEROL		127	Desirable: Less than 130 Above Desirable: 130 - 159 mg/dL

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			Borderline High: 160 - 189 High: 190 - 219 Very high: > or = 220
METHOD : CALCULATED PARAMETER			
CHOL/HDL RATIO		3.3	3.3 - 4.4 Low Risk 4.5 - 7.0 Average Risk 7.1 - 11.0 Moderate Risk > 11.0 High Risk
METHOD : CALCULATED PARAMETER			
LDL/HDL RATIO		1.9	0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk
METHOD : CALCULATED PARAMETER			
VERY LOW DENSITY LIPOPROTEIN		23.6	</= 30.0 mg/dL
METHOD : CALCULATED PARAMETER			

Interpretation(s)

LIVER FUNCTION PROFILE, SERUM- LIVER FUNCTION PROFILE

Bilirubin is a yellowish pigment found in bile and is a breakdown product of normal heme catabolism. Bilirubin is excreted in bile and urine, and elevated levels may give yellow discoloration in jaundice. Elevated levels result from increased bilirubin production (eg, hemolysis and ineffective erythropoiesis), decreased bilirubin excretion (eg, obstruction and hepatitis), and abnormal bilirubin metabolism (eg, hereditary and neonatal jaundice). Conjugated (direct) bilirubin is elevated more than unconjugated (indirect) bilirubin in Viral hepatitis, Drug reactions, Alcoholic liver disease Conjugated (direct) bilirubin is also elevated more than unconjugated (indirect) bilirubin when there is some kind of blockage of the bile ducts like in Gallstones getting into the bile ducts, tumors & Scarring of the bile ducts. Increased unconjugated (indirect) bilirubin may be a result of Hemolytic or pernicious anemia, Transfusion reaction & a common metabolic condition termed Gilbert syndrome, due to low levels of the enzyme that attaches sugar molecules to bilirubin.

AST is an enzyme found in various parts of the body. AST is found in the liver, heart, skeletal muscle, kidneys, brain, and red blood cells, and it is commonly measured clinically as a marker for liver health. AST levels increase during chronic viral hepatitis, blockage of the bile duct, cirrhosis of the liver, liver cancer, kidney failure, hemolytic anemia, pancreatitis, hemochromatosis. AST levels may also increase after a heart attack or strenuous activity. ALT test measures the amount of this enzyme in the blood. ALT is found mainly in the liver, but also in smaller amounts in the kidneys, heart, muscles, and pancreas. It is commonly measured as a part of a diagnostic evaluation of hepatocellular injury, to determine liver health. AST levels increase during acute hepatitis, sometimes due to a viral infection, ischemia to the liver, chronic hepatitis, obstruction of bile ducts, cirrhosis.

ALP is a protein found in almost all body tissues. Tissues with higher amounts of ALP include the liver, bile ducts and bone. Elevated ALP levels are seen in Biliary obstruction, Osteoblastic bone tumors, osteomalacia, hepatitis, Hyperparathyroidism, Leukemia, Lymphoma, Paget's disease, Rickets, Sarcoidosis etc. Lower-than-normal ALP levels seen in Hypophosphatasia, Malnutrition, Protein deficiency, Wilson's disease. GGT is an enzyme found in cell membranes of many tissues mainly in the liver, kidney and pancreas. It is also found in other tissues including intestine, spleen, heart, brain and seminal vesicles. The highest concentration is in the kidney, but the liver is considered the source of normal enzyme activity. Serum GGT has been widely used as an index of liver dysfunction. Elevated serum GGT activity can be found in diseases of the liver, biliary system and pancreas. Conditions that increase serum GGT are obstructive liver disease, high alcohol consumption and use of enzyme-inducing drugs etc. Serum total protein, also known as total protein, is a biochemical test for measuring the total amount of protein in serum. Protein in the plasma is made up of albumin and globulin. Higher-than-normal levels may be due to: Chronic inflammation or infection, including HIV and hepatitis B or C, Multiple myeloma, Waldenstrom's disease. Lower-than-normal levels may be due to: Agammaglobulinemia, Bleeding (hemorrhage), Burns, Glomerulonephritis, Liver disease, Malabsorption, Malnutrition, Nephrotic syndrome, Protein-losing enteropathy etc. Human serum albumin is the most abundant protein in human blood plasma. It is produced in the liver. Albumin constitutes about half of the blood serum protein. Low blood albumin levels (hypoalbuminemia) can be caused by: Liver disease like cirrhosis of the liver, nephrotic syndrome, protein-losing enteropathy, Burns, hemodilution, increased vascular permeability or decreased lymphatic clearance, malnutrition and wasting etc

GLUCOSE, FASTING, PLASMA-

ADA 2021 guidelines for adults, after 8 hrs fasting is as follows:

Pre-diabetics: 100 - 125 mg/dL

Diabetic: > or = 126 mg/dL

GLYCOSYLATED HEMOGLOBIN, EDTA WHOLE BLOOD-

Glycosylated hemoglobin (GHb) has been firmly established as an index of long-term blood glucose concentrations and as a measure of the risk for the development of complications in patients with diabetes mellitus. Formation of GHb is essentially irreversible, and the concentration in the blood depends on both the life span of the red blood cell (average 120 days) and the blood glucose concentration. Because the rate of formation of GHb is directly proportional to the concentration of glucose in the blood, the GHb concentration represents the integrated values for glucose over the preceding 6-8 weeks.

Any condition that alters the life span of the red blood cells has the potential to alter the GHb level. Samples from patients with hemolytic anemias will exhibit decreased glycated hemoglobin values due to the shortened life span of the red cells. This effect will depend upon the severity of the anemia. Samples from patients with polycythemia or post-splenectomy may exhibit increased glycated hemoglobin values due to a somewhat longer life span of the red cells.

Glycosylated hemoglobins results from patients with HbSS, HbCC, and HbSC and HbD must be interpreted with caution, given the pathological processes, including anemia, increased red cell turnover, transfusion requirements, that adversely impact HbA1c as a marker of long-term glycemic control. In these conditions, alternative forms of

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**PATIENT NAME : MS.PINKI KUMARI RAJAK**PATIENT ID : **FH.12051116**

CLIENT PATIENT ID : UID:12051116

ACCESSION NO : **0022VJ001489** AGE : 40 Years SEX : Female DATE OF BIRTH : 17/06/1982

DRAWN : 08/10/2022 10:10 RECEIVED : 08/10/2022 10:10 REPORTED : 08/10/2022 14:59

CLIENT NAME : **FORTIS VASHI-CHC -SPLZD**

REFERRING DOCTOR : SELF

CLINICAL INFORMATION :

UID:12051116 REQNO-1305061

CORP-OPD

BILLNO-150122OPCR050212

BILLNO-150122OPCR050212

Test Report Status	Final	Results	Biological Reference Interval
--------------------	-------	---------	-------------------------------

testing such as glycated serum protein (fructosamine) should be considered.

"Targets should be individualized; More or less stringent glycemic goals may be appropriate for individual patients. Goals should be individualized based on duration of diabetes, age/life expectancy, comorbid conditions, known CVD or advanced microvascular complications, hypoglycemia unawareness, and individual patient considerations."

References

1. Tietz Textbook of Clinical Chemistry and Molecular Diagnostics, edited by Carl A Burtis, Edward R.Ashwood, David E Bruns, 4th Edition, Elsevier publication, 2006, 879-884.
 2. Forsham PH. Diabetes Mellitus:A rational plan for management. Postgrad Med 1982, 71,139-154.
 3. Mayer TK, Freedman ZR: Protein glycosylation in Diabetes Mellitus: A review of laboratory measurements and their clinical utility. Clin Chim Acta 1983, 127, 147-184.
- CORONARY RISK PROFILE (LIPID PROFILE), SERUM-Serum cholesterol** is a blood test that can provide valuable information for the risk of coronary artery disease This test can help determine your risk of the build up of plaques in your arteries that can lead to narrowed or blocked arteries throughout your body (atherosclerosis). High cholesterol levels usually don't cause any signs or symptoms, so a cholesterol test is an important tool. High cholesterol levels often are a significant risk factor for heart disease and important for diagnosis of hyperlipoproteinemia, atherosclerosis, hepatic and thyroid diseases.

Serum Triglyceride are a type of fat in the blood. When you eat, your body converts any calories it doesn't need into triglycerides, which are stored in fat cells. High triglyceride levels are associated with several factors, including being overweight, eating too many sweets or drinking too much alcohol, smoking, being sedentary, or having diabetes with elevated blood sugar levels. Analysis has proven useful in the diagnosis and treatment of patients with diabetes mellitus, nephrosis, liver obstruction, other diseases involving lipid metabolism, and various endocrine disorders. In conjunction with high density lipoprotein and total serum cholesterol, a triglyceride determination provides valuable information for the assessment of coronary heart disease risk.It is done in fasting state.

High-density lipoprotein (HDL) cholesterol. This is sometimes called the "good" cholesterol because it helps carry away LDL cholesterol, thus keeping arteries open and blood flowing more freely.HDL cholesterol is inversely related to the risk for cardiovascular disease. It increases following regular exercise, moderate alcohol consumption and with oral estrogen therapy. Decreased levels are associated with obesity, stress, cigarette smoking and diabetes mellitus.

SERUM LDL The small dense LDL test can be used to determine cardiovascular risk in individuals with metabolic syndrome or established/progressing coronary artery disease, individuals with triglyceride levels between 70 and 140 mg/dL, as well as individuals with a diet high in trans-fat or carbohydrates. Elevated sdLDL levels are associated with metabolic syndrome and an 'atherogenic lipoprotein profile', and are a strong, independent predictor of cardiovascular disease. Elevated levels of LDL arise from multiple sources. A major factor is sedentary lifestyle with a diet high in saturated fat. Insulin-resistance and pre-diabetes have also been implicated, as has genetic predisposition. Measurement of sdLDL allows the clinician to get a more comprehensive picture of lipid risk factors and tailor treatment accordingly. Reducing LDL levels will reduce the risk of CVD and MI.

Non HDL Cholesterol - Adult treatment panel ATP III suggested the addition of Non-HDL Cholesterol as an indicator of all atherogenic lipoproteins (mainly LDL and VLDL). NICE guidelines recommend Non-HDL Cholesterol measurement before initiating lipid lowering therapy. It has also been shown to be a better marker of risk in both primary and secondary prevention studies.

Recommendations:

Results of Lipids should always be interpreted in conjunction with the patient's medical history, clinical presentation and other findings.

NON FASTING LIPID PROFILE includes Total Cholesterol, HDL Cholesterol and calculated non-HDL Cholesterol. It does not include triglycerides and may be best used in patients for whom fasting is difficult.

CLINICAL PATH**URINALYSIS****PHYSICAL EXAMINATION, URINE**

COLOR PALE YELLOW

METHOD : PHYSICAL

APPEARANCE SLIGHTLY HAZY

METHOD : VISUAL

SPECIFIC GRAVITY 1.015 1.003 - 1.035

METHOD : REFLECTANCE SPECTROPHOTOMETRY (APPARENT PKA CHANGE OF PRETREATED POLYELECTROLYTES IN RELATION TO IONIC CONCENTRATION)

CHEMICAL EXAMINATION, URINE

PH 6.0 4.7 - 7.5

METHOD : REFLECTANCE SPECTROPHOTOMETRY- DOUBLE INDICATOR METHOD

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PATIENT NAME : MS.PINKI KUMARI RAJAKPATIENT ID : **FH.12051116**

CLIENT PATIENT ID : UID:12051116

ACCESSION NO : **0022VJ001489** AGE : 40 Years SEX : Female DATE OF BIRTH : 17/06/1982

DRAWN : 08/10/2022 10:10

RECEIVED : 08/10/2022 10:10

REPORTED : 08/10/2022 14:59

CLIENT NAME : **FORTIS VASHI-CHC -SPLZD**

REFERRING DOCTOR : SELF

CLINICAL INFORMATION :

UID:12051116 REQNO-1305061

CORP-OPD

BILLNO-150122OPCR050212

BILLNO-150122OPCR050212

Test Report Status	Final	Results	Biological Reference Interval
PROTEIN		NOT DETECTED	NOT DETECTED
METHOD : REFLECTANCE SPECTROPHOTOMETRY - PROTEIN-ERROR-OF-INDICATOR PRINCIPLE			
GLUCOSE		NOT DETECTED	NOT DETECTED
METHOD : REFLECTANCE SPECTROPHOTOMETRY, DOUBLE SEQUENTIAL ENZYME REACTION-GOD/POD			
KETONES		NOT DETECTED	NOT DETECTED
METHOD : REFLECTANCE SPECTROPHOTOMETRY, ROTHERA'S PRINCIPLE			
BLOOD		DETECTED (TRACE) IN URINE	
METHOD : REFLECTANCE SPECTROPHOTOMETRY, PEROXIDASE LIKE ACTIVITY OF HAEMOGLOBIN			
BILIRUBIN		NOT DETECTED	NOT DETECTED
METHOD : REFLECTANCE SPECTROPHOTOMETRY, DIAZOTIZATION- COUPLING OF BILIRUBIN WITH DIAZOTIZED SALT			
UROBILINOGEN		NORMAL	NORMAL
METHOD : REFLECTANCE SPECTROPHOTOMETRY (MODIFIED EHRlich REACTION)			
NITRITE		NOT DETECTED	NOT DETECTED
METHOD : REFLECTANCE SPECTROPHOTOMETRY, CONVERSION OF NITRATE TO NITRITE			
LEUKOCYTE ESTERASE		NOT DETECTED	NOT DETECTED
METHOD : REFLECTANCE SPECTROPHOTOMETRY, ESTERASE HYDROLYSIS ACTIVITY			
MICROSCOPIC EXAMINATION, URINE			
PUS CELL (WBC'S)		3-5	0-5 /HPF
METHOD : MICROSCOPIC EXAMINATION			
EPITHELIAL CELLS		15-20	0-5 /HPF
METHOD : MICROSCOPIC EXAMINATION			
ERYTHROCYTES (RBC'S)		1 - 2	NOT DETECTED /HPF
METHOD : MICROSCOPIC EXAMINATION			
CASTS		NOT DETECTED	
METHOD : MICROSCOPIC EXAMINATION			
CRYSTALS		NOT DETECTED	
METHOD : MICROSCOPIC EXAMINATION			
BACTERIA		DETECTED	NOT DETECTED
METHOD : MICROSCOPIC EXAMINATION			
YEAST		NOT DETECTED	NOT DETECTED
METHOD : MICROSCOPIC EXAMINATION			
REMARKS		URINARY MICROSCOPIC EXAMINATION DONE ON URINARY CENTRIFUGED SEDIMENT	

Interpretation(s)**MICROSCOPIC EXAMINATION, URINE-**

Routine urine analysis assists in screening and diagnosis of various metabolic, urological, kidney and liver disorders

Protein: Elevated proteins can be an early sign of kidney disease. Urinary protein excretion can also be temporarily elevated by strenuous exercise, orthostatic proteinuria, dehydration, urinary tract infections and acute illness with fever

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**PATIENT NAME : MS.PINKI KUMARI RAJAK**PATIENT ID : **FH.12051116**

CLIENT PATIENT ID : UID:12051116

ACCESSION NO : **0022VJ001489** AGE : 40 Years SEX : Female DATE OF BIRTH : 17/06/1982

DRAWN : 08/10/2022 10:10 RECEIVED : 08/10/2022 10:10 REPORTED : 08/10/2022 14:59

CLIENT NAME : **FORTIS VASHI-CHC -SPLZD**

REFERRING DOCTOR : SELF

CLINICAL INFORMATION :

UID:12051116 REQNO-1305061

CORP-OPD

BILLNO-150122OPCR050212

BILLNO-150122OPCR050212

Test Report Status	Final	Results	Biological Reference Interval
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Glucose: Uncontrolled diabetes mellitus can lead to presence of glucose in urine. Other causes include pregnancy, hormonal disturbances, liver disease and certain medications.

Ketones: Uncontrolled diabetes mellitus can lead to presence of ketones in urine. Ketones can also be seen in starvation, frequent vomiting, pregnancy and strenuous exercise.

Blood: Occult blood can occur in urine as intact erythrocytes or haemoglobin, which can occur in various urological, nephrological and bleeding disorders.

Leukocytes: An increase in leukocytes is an indication of inflammation in urinary tract or kidneys. Most common cause is bacterial urinary tract infection.

Nitrite: Many bacteria give positive results when their number is high. Nitrite concentration during infection increases with length of time the urine specimen is retained in bladder prior to collection.

pH: The kidneys play an important role in maintaining acid base balance of the body. Conditions of the body producing acidosis/ alkalosis or ingestion of certain type of food can affect the pH of urine.

Specific gravity: Specific gravity gives an indication of how concentrated the urine is. Increased specific gravity is seen in conditions like dehydration, glycosuria and proteinuria while decreased specific gravity is seen in excessive fluid intake, renal failure and diabetes insipidus.

Bilirubin: In certain liver diseases such as biliary obstruction or hepatitis, bilirubin gets excreted in urine.

Urobilinogen: Positive results are seen in liver diseases like hepatitis and cirrhosis and in cases of hemolytic anemia

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Dr. Rekha Nair, MD
Microbiologist

Dr. Akta Dubey
Consultant Pathologist

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Patient Ref. No. 2200000800581

**PATIENT NAME : MS.PINKI KUMARI RAJAK**PATIENT ID : **FH.12051116**

CLIENT PATIENT ID : UID:12051116

ACCESSION NO : **0022VJ001489** AGE : 40 Years SEX : Female DATE OF BIRTH : 17/06/1982

DRAWN : 08/10/2022 10:10

RECEIVED : 08/10/2022 10:10

REPORTED : 08/10/2022 17:11

CLIENT NAME : **FORTIS VASHI-CHC -SPLZD**

REFERRING DOCTOR : SELF

CLINICAL INFORMATION :

UID:12051116 REQNO-1305061

CORP-OPD

BILLNO-150122OPCR050212

BILLNO-150122OPCR050212

Test Report Status	Final	Results	Biological Reference Interval	Units
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SPECIALISED CHEMISTRY - HORMONE**THYROID PANEL, SERUM**

T3	155.5	80 - 200	ng/dL
METHOD : ELECTROCHEMILUMINESCENCE, COMPETITIVE IMMUNOASSAY			
T4	10.13	5.1 - 14.1	µg/dL
METHOD : ELECTROCHEMILUMINESCENCE, COMPETITIVE IMMUNOASSAY			
TSH 3RD GENERATION	4.240	High 0.270 - 4.200	µIU/mL
METHOD : ELECTROCHEMILUMINESCENCE, COMPETITIVE IMMUNOASSAY			

Comments

NOTE: PLEASE CORRELATE VALUES OF THYROID FUNCTION TEST WITH THE CLINICAL & TREATMENT HISTORY OF THE PATIENT.

Interpretation(s)**THYROID PANEL, SERUM-**

Triiodothyronine T3, is a thyroid hormone. It affects almost every physiological process in the body, including growth, development, metabolism, body temperature, and heart rate. Production of T3 and its prohormone thyroxine (T4) is activated by thyroid-stimulating hormone (TSH), which is released from the pituitary gland. Elevated concentrations of T3, and T4 in the blood inhibit the production of TSH.

Thyroxine T4, Thyroxine's principal function is to stimulate the metabolism of all cells and tissues in the body. Excessive secretion of thyroxine in the body is hyperthyroidism, and deficient secretion is called hypothyroidism. Most of the thyroid hormone in blood is bound to transport proteins. Only a very small fraction of the circulating hormone is free and biologically active.

In primary hypothyroidism, TSH levels are significantly elevated, while in secondary and tertiary hypothyroidism, TSH levels are low.

Below mentioned are the guidelines for Pregnancy related reference ranges for Total T4, TSH & Total T3

Levels in	TOTAL T4 (µg/dL)	TSH3G (µIU/mL)	TOTAL T3 (ng/dL)
Pregnancy	6.6 - 12.4	0.1 - 2.5	81 - 190
1st Trimester	6.6 - 12.4	0.1 - 2.5	81 - 190
2nd Trimester	6.6 - 15.5	0.2 - 3.0	100 - 260
3rd Trimester	6.6 - 15.5	0.3 - 3.0	100 - 260

Below mentioned are the guidelines for age related reference ranges for T3 and T4.

	T3 (ng/dL)	T4 (µg/dL)
New Born:	75 - 260	1-3 day: 8.2 - 19.9
		1 Week: 6.0 - 15.9

NOTE: TSH concentrations in apparently normal euthyroid subjects are known to be highly skewed, with a strong tailed distribution towards higher TSH values. This is well documented in the pediatric population including the infant age group.

Kindly note: Method specific reference ranges are appearing on the report under biological reference range.

Reference:

- Burtis C.A., Ashwood E. R. Bruns D.E. Teitz textbook of Clinical Chemistry and Molecular Diagnostics, 4th Edition.
- Gowenlock A.H. Varley's Practical Clinical Biochemistry, 6th Edition.
- Behrman R.E. Kliegman R.M., Jenson H. B. Nelson Text Book of Pediatrics, 17th Edition

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Patient Ref. No. 22000000800581

**PATIENT NAME : MS.PINKI KUMARI RAJAK**PATIENT ID : **FH.12051116**

CLIENT PATIENT ID : UID:12051116

ACCESSION NO : **0022VJ001489**

AGE : 40 Years

SEX : Female

DATE OF BIRTH : 17/06/1982

DRAWN : 08/10/2022 10:10

RECEIVED : 08/10/2022 10:10

REPORTED : 08/10/2022 17:11

CLIENT NAME : **FORTIS VASHI-CHC -SPLZD**

REFERRING DOCTOR : SELF

CLINICAL INFORMATION :

UID:12051116 REQNO-1305061

CORP-OPD

BILLNO-150122OPCR050212

BILLNO-150122OPCR050212

Test Report Status	Results	Biological Reference Interval	Units
Final			

Dr. Swapnil Sirmukaddam
Consultant Pathologist

SRL Ltd

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**PATIENT NAME : MS.PINKI KUMARI RAJAK**PATIENT ID : **FH.12051116**

CLIENT PATIENT ID : UID:12051116

ACCESSION NO : **0022VJ001552** AGE : 40 Years SEX : Female DATE OF BIRTH : 17/06/1982

DRAWN : 08/10/2022 12:53

RECEIVED : 08/10/2022 12:53

REPORTED : 08/10/2022 13:57

CLIENT NAME : **FORTIS VASHI-CHC -SPLZD**

REFERRING DOCTOR :

CLINICAL INFORMATION :

UID:12051116 REQNO-1305061

CORP-OPD

BILLNO-150122OPCR050212

BILLNO-150122OPCR050212

Test Report Status	Final	Results	Biological Reference Interval	Units
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BIO CHEMISTRY**GLUCOSE, POST-PRANDIAL, PLASMA**

GLUCOSE, POST-PRANDIAL, PLASMA

141**High** 70 - 139

mg/dL

METHOD : HEXOKINASE

Interpretation(s)

GLUCOSE, POST-PRANDIAL, PLASMA-ADA Guidelines for 2hr post prandial glucose levels is only after ingestion of 75grams of glucose in 300 ml water, over a period of 5 minutes.

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Dr.Akta Dubey**Consultant Pathologist****SRL Ltd**

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**Patient Ref. No. 22000000800644**

PATIENT NAME : MS.PINKI KUMARI RAJAKPATIENT ID : **FH.12051116**

CLIENT PATIENT ID : UID:12051116

ACCESSION NO : **0022VJ001595** AGE : 40 Years SEX : Female DATE OF BIRTH : 17/06/1982

DRAWN : 08/10/2022 14:09 RECEIVED : 08/10/2022 14:29 REPORTED : 10/10/2022 10:35

CLIENT NAME : **FORTIS VASHI-CHC -SPLZD**

REFERRING DOCTOR :

CLINICAL INFORMATION :

UID:12051116 REQNO-1305061

CORP-OPD

BILLNO-1501220PCR050212

BILLNO-1501220PCR050212

Test Report Status

Final

Units

CYTOLOGY**PAPANICOLAOU SMEAR****PAPANICOLAOU SMEAR**

TEST METHOD

CONVENTIONAL GYNEC CYTOLOGY

SPECIMEN TYPE

TWO UNSTAINED CERVICAL SMEARS RECEIVED

REPORTING SYSTEM

2014 BETHESDA SYSTEM FOR REPORTING CERVICAL CYTOLOGY

SPECIMEN ADEQUACY

SATISFACTORY

METHOD : MICROSCOPIC EXAMINATION

MICROSCOPY

SMEARS STUDIED SHOW SUPERFICIAL SQUAMOUS CELLS, INTERMEDIATE SQUAMOUS CELLS, OCCASIONAL SQUAMOUS METAPLASTIC CELLS, OCCASIONAL CLUSTERS OF ENDOCERVICAL CELLS IN THE BACKGROUND OF FEW POLYMORPHS.

INTERPRETATION / RESULT

NEGATIVE FOR INTRAEPITHELIAL LESION OR MALIGNANCY

Comments

PLEASE NOTE PAPANICOLAOU SMEAR STUDY IS A SCREENING PROCEDURE FOR CERVICAL CANCER WITH INHERENT FALSE NEGATIVE RESULTS, HENCE SHOULD BE INTERPRETED WITH CAUTION.

NO CYTOLOGICAL EVIDENCE OF HPV INFECTION IN THE SMEARS STUDIED.

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Dr.Akta Dubey

Consultant Pathologist

SRL Ltd

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Patient Ref. No. 22000000800687

40 Years

Female

HC

NSR

Rate 60 . Sinus rhythm.....normal P axis, V-rate 50-99

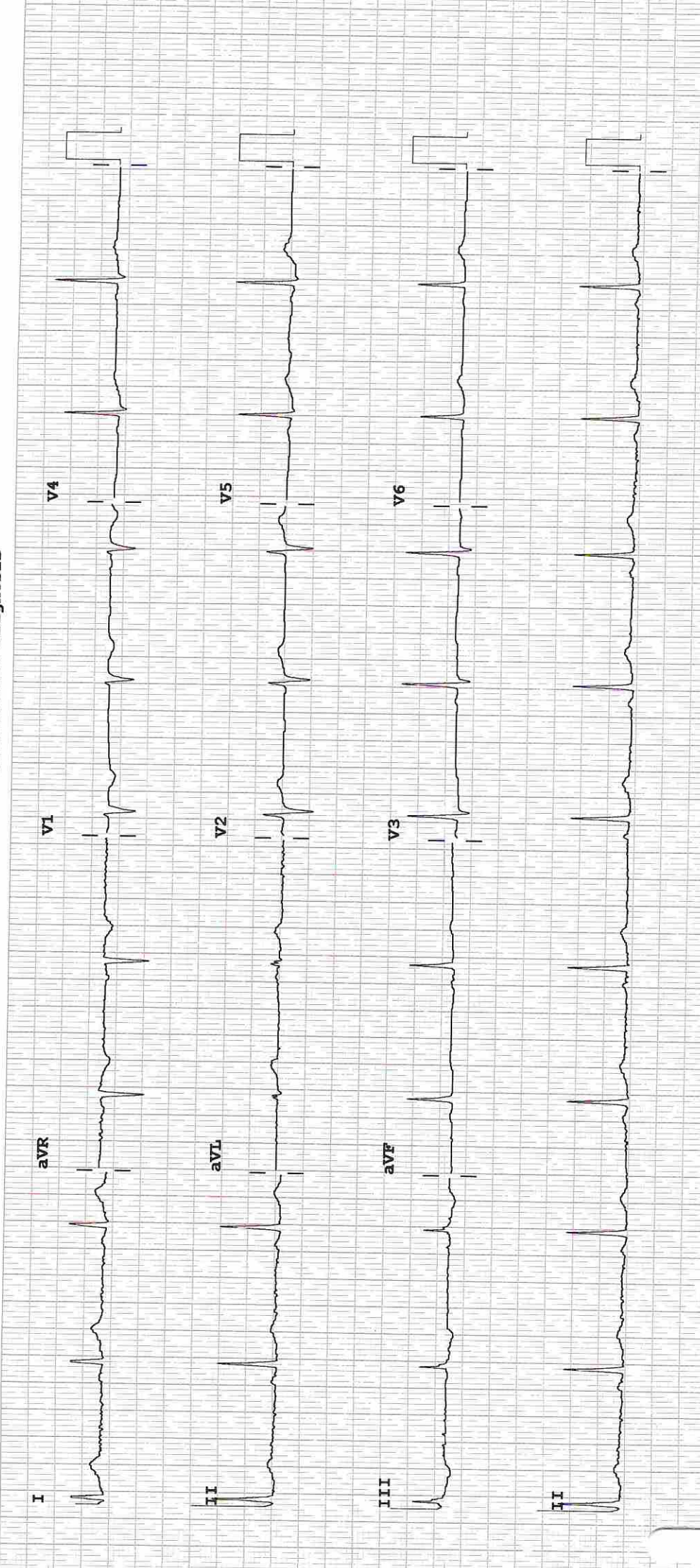
PR 134
QRS 83
QT 370
QTc 370

--AXIS--
P 33
QRS 53
T 14

12 Lead; Standard Placement

- NORMAL ECG -

Unconfirmed Diagnosis



Device: Speed: 25 mm/sec Limb: 10 mm/mV Chest: 10.0 mm/mV

F 50~ 0.50-100 Hz W

100B CL P?



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DEPARTMENT OF NIC

Date: 08/Oct/2022

Name: Ms. Pinki Kumari Rajak

UHID | Episode No : 12051116 | 49889/22/1501

Age | Sex: 40 YEAR(S) | Female

Order No | Order Date: 1501/PN/OP/2210/105565 | 08-Oct-2022

Order Station : FO-OPD

Admitted On | Reporting Date : 08-Oct-2022 16:34:23

Bed Name :

Order Doctor Name : Dr.SELF .

ECHOCARDIOGRAPHY TRANSTHORACIC

FINDINGS:

- No left ventricle regional wall motion abnormality at rest.
- Normal left ventricle systolic function. LVEF = 60%.
- No left ventricle diastolic dysfunction.
- No mitral regurgitation.
- No aortic regurgitation. No aortic stenosis.
- Trivial tricuspid regurgitation. No pulmonary hypertension. PASP = 25 mm of Hg.
- Intact IVS and IAS.
- No left ventricle clot/vegetation/pericardial effusion.
- Normal right atrium and right ventricle dimension.
- Normal left atrium and left ventricle dimension.
- Normal right ventricle systolic function. No hepatic congestion.

M-MODE MEASUREMENTS:

LA	35	mm
AO Root	29	mm
AO CUSP SEP	18	mm
LVID (s)	31	mm
LVID (d)	43	mm
IVS (d)	10	mm
LVPW (d)	09	mm
RVID (d)	29	mm
RA	31	mm
LVEF	60	%



(For Billing/Reports & Discharge Summary only)

DEPARTMENT OF NIC

Date: 08/Oct/2022

Name: Ms. Pinki Kumari Rajak
Age | Sex: 40 YEAR(S) | Female
Order Station : FO-OPD
Bed Name :

UHID | Episode No : 12051116 | 49889/22/1501
Order No | Order Date: 1501/PN/OP/2210/105565 | 08-Oct-2022
Admitted On | Reporting Date : 08-Oct-2022 16:34:23
Order Doctor Name : Dr.SELF .

DOPPLER STUDY:

E WAVE VELOCITY: 0.7 m/sec.
A WAVE VELOCITY: 0.8 m/sec
E/A RATIO: 0.6

	PEAK (mmHg)	MEAN (mmHg)	V max (m/sec)	GRADE OF REGURGITATION
MITRAL VALVE	N			Nil
AORTIC VALVE	05			Nil
TRICUSPID VALVE	25			Trivial
PULMONARY VALVE	2.0			Nil

Final Impression :

- No RWMA.
- Trivial TR. No PH.
- Normal LV and RV systolic function.


DR. PRASHANT PAWAR,
DNB(MED), DNB(CARDIOLOGY)



(For Billing/Reports & Discharge Summary only)

DEPARTMENT OF RADIOLOGY

Date: 10/Oct/2022

Name: Ms. Pinki Kumari Rajak

UHID | Episode No : 12051116 | 49889/22/1501

Age | Sex: 40 YEAR(S) | Female

Order No | Order Date: 1501/PN/OP/2210/105565 | 08-Oct-2022

Order Station : FO-OPD

Admitted On | Reporting Date : 10-Oct-2022 15:31:44

Bed Name :

Order Doctor Name : Dr.SELF .

X-RAY-CHEST- PA

Findings:

Both lung fields are clear.

The cardiac shadow appears within normal limits.

Trachea and major bronchi appears normal.

Both costophrenic angles are well maintained.

Bony thorax is unremarkable.

DR. ABHIJEET BHAMBURE
DMRD, DNB (Radiologist)



(For Billing/Reports & Discharge Summary only)

DEPARTMENT OF RADIOLOGY

Date: 08/Oct/2022

Name: Ms. Pinki Kumari Rajak

Age | Sex: 40 YEAR(S) | Female

Order Station : FO-OPD

Bed Name :

UHID | Episode No : 12051116 | 49889/22/1501

Order No | Order Date: 1501/PN/OP/2210/105565 | 08-Oct-2022

Admitted On | Reporting Date : 08-Oct-2022 15:14:15

Order Doctor Name : Dr.SELF .

US-WHOLE ABDOMEN (TAS)

LIVER is normal in size (14.8 cm) and echogenicity. Intrahepatic portal and biliary systems are normal. No focal lesion is seen in liver. Portal vein is normal.

GALL BLADDER is physiologically distended. Gall bladder reveals normal wall thickness. No evidence of calculi in gall bladder. No evidence of pericholecystic collection.

SPLEEN is normal in size (9.5 cm) and echogenicity.

BOTH KIDNEYS are normal in size and echogenicity. The central sinus complex is normal. No evidence of calculi/hydronephrosis.

Right kidney measures 9.5 x 4.1 cm.

Left kidney measures 8.3 x 3.8 cm.

PANCREAS is normal in size and morphology. No evidence of peripancreatic collection.

URINARY BLADDER is normal in capacity and contour. Bladder wall is normal in thickness. No evidence of intravesical mass/calculi.

UTERUS is normal in size, measuring 7.4 x 4.2 x 5.3 cm.
Endometrium measures 7.0 mm in thickness.

Right ovary is normal.

Left ovary is not visualised, however the adnexa is clear.

No evidence of ascites.

Impression:

- No significant abnormality is detected.

DR. YOGESH PATHADE
(MD Radio-diagnosis)



(For Billing/Reports & Discharge Summary only)

DEPARTMENT OF RADIOLOGY

Date: 08/Oct/2022

Name: Ms. Pinki Kumari Rajak
Age | Sex: 40 YEAR(S) | Female
Order Station : FO-OPD
Bed Name :

UHID | Episode No : 12051116 | 49889/22/1501
Order No | Order Date: 1501/PN/OP/2210/105565 | 08-Oct-2022
Admitted On | Reporting Date : 08-Oct-2022 16:45:03
Order Doctor Name : Dr.SELF .

MAMMOGRAM - BOTH BREAST

Findings:

Bilateral film screen mammography was performed in cranio-caudal and medio-lateral oblique views.

Both breasts show scattered areas of fibroglandular density.

No evidence of any dominant mass, clusters of microcalcifications, nipple retraction, skin thickening or abnormal vascularity is seen in either breast.

No evidence of axillary lymphadenopathy.

IMPRESSION:

- No significant abnormality detected. (BI-RADS category I).
- No obvious mass lesion in the breasts.

Normal-interval follow-up is recommended.

DR. YOGINI SHAH
DMRD., DNB. (Radiologist)